AMENDMENT UNDER 37 C.F.R. § 1.114(c) Attorney Docket No.: A8709

U.S. Application No.: 10/823,647

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1.-26. (canceled).

(currently amended): A method for diagnosing a bipolar disorder in a <u>human</u>

patient, comprising:

- (a) obtaining a patient ratio of
 - the mean membrane potential of fresh-cells of the a test human patient incubated in vitro in the presence of a compound that alters Na⁺K⁺
 ATPase activity, but in the absence of K⁺, to
 - (ii) the mean membrane potential of fresh-cells of the test human patient incubated in vitro in the absence of the compound that alters Na⁺K⁺
 ATPase activity, but in the presence of K⁺; and

one or both of the following steps (b) and (c):

- (b) comparing the patient-ratio obtained in (a) to a control ratio, wherein the control ratio is the ratio of
 - (iii) the mean membrane potential of corresponding fresh-control cells of one or more people-humans known to not have said bipolar disorder incubated in vitro in the presence of a compound that alters Na⁺K⁺ ATPase activity, but in the absence of K⁺, to

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(iv) the mean membrane potential of corresponding fresh-control cells of one or more people-humans known to not have said bipolar disorder incubated in vitro in the absence of the compound that alters Na⁺K⁺ ATPase activity, but in the presence of K⁺.

wherein awhen, significantly lower patient the ratio obtained in (a) is

significantly lower than compared to the control ratio obtained in (b), said

indicates that the test human patient is diagnosed as having has said bipolar
disorder:

- (c) comparing the patient-ratio obtained in (a) to a bipolar control ratio, wherein the bipolar control ratio is the ratio of
 - (v) the mean membrane potential of corresponding fresh-bipolar control cells of one or more people-humans known to have said bipolar disorder incubated in vitro in the presence of a compound that alters Na⁺K⁺ ATPase activity, but in the absence of K⁺₋, to
 - (vi) the mean membrane potential of corresponding fresh-bipolar control cells of one or more people-humans known to have said bipolar disorder incubated in vitro in the absence of the compound that alters Na⁺K⁺ ATPase activity, but in the presence of K⁺,
 - wherein when the lack of a significant difference between the patient-ratio

 obtained in (a) is not significantly different than compared to the bipolar control ratio obtained in (c), said indicates that the test human patient is

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diagnosed as havinghas said_ bipolar disorder, wherein said patient is a

human_;

wherein the cells incubated in vitro in the presence of the compound that alters Na+K+

ATPase activity are incubated in vitro in the absence of K+, and wherein the cells incubated

in vitro in the absence of the compound that alters Na K-ATPase activity are incubated in vitro

in the presence of K+

wherein each mean membrane potential is determined by incubating the cells in vitro in

buffer comprising a potential-sensitive dye, resuspending the cells in potential-sensitive dye

free-buffer, and measuring cell fluorescence.

28.-30. (canceled).

31. (currently amended): The method according to claim 27, wherein the compound

that alters Na⁺K⁺ ATPase activity is selected from the group consisting of: valinomycin,

monensin, monensin decyl ester, gramieidin, p-chloromercurybenzenesulfonate (PCMBS).

veratridine, ethacrynate, dopamine, a catecholamine, a phorbol ester, ouabain, lithium, valproate,

lamotrigine, cocaine, nicotine, R0-31-8220, oxymetazoline, calcineurin, topiramate, a peptide

hormone, sorbitol, and a diuretic.

32. (original) The method according to claim 31, wherein the compound that alters

Na+K+ ATPase activity is ethacrynate.

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33.-44. (canceled).

45. (previously presented) The method of claim 27, wherein said bipolar disorder is bipolar I disorder.

46.-51. (canceled).